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| APPLICATION NO. | FILING DATE | FIRST NAMED INVENTOR | ATTORNEY DOCKET NO. | CONFIRMATION NO. | |
|--------------------------------|----------------------|----------------------|---------------------------------|------------------|--|
| 09/936,985 | 12/19/2001 | Carine Capiau | B45182 | 2966 | |
| | 7590 02/21/2002 | | | | |
| SMITHKLINE BEECHAM CORPORATION | | | ЕХАМП | EXAMINER | |
| P. O. BOX 15 | P. O. BOX 1539 | | | FORD, VANESSA L | |
| KING OF PR | USSIA, PA 19406-0939 | | ART UNIT | PAPER NUMBER | |
| | | | 1645 DATE MAILED: 02/21/2002 | 6 | |

Please find below and/or attached an Office communication concerning this application or proceeding.

| \ 4 | | \downarrow | ale | | | |
|---|--|--|--|--|--|--|
| | | Application No. | Applicant(s) | | | |
| Office Action Summary | | 09/936,985 | CAPIAU ET AL. | | | |
| | | Examiner | Art Unit | | | |
| | | Vanessa L. Ford | 1645 | | | |
| Period fo | The MAILING DATE of this communication apport Reply | pears on the cover sheet wi | th the correspondence address | | | |
| THE - Externanter - If the - If NC - Failu - Any | ORTENED STATUTORY PERIOD FOR REPLY MAILING DATE OF THIS COMMUNICATION. Insions of time may be available under the provisions of 37 CFR 1.1 SIX (6) MONTHS from the mailing date of this communication. In period for reply specified above is less than thirty (30) days, a reply period for reply is specified above, the maximum statutory period for reply within the set or extended period for reply will, by statute reply received by the Office later than three months after the mailing and patent term adjustment. See 37 CFR 1.704(b). | 36(a). In no event, however, may a re y within the statutory minimum of thirty will apply and will expire SIX (6) MON' , cause the application to become AB | eply be timely filed (30) days will be considered timely. FHS from the mailing date of this communication. ANDONED (35 U.S.C. § 133). | | | |
| 1)🖂 | Responsive to communication(s) filed on 195 | September 2001 . | | | | |
| 2a)□ | This action is FINAL . 2b) ☐ Th | is action is non-final. | | | | |
| 3)□ | 3) Since this application is in condition for allowance except for formal matters, prosecution as to the merits is closed in accordance with the practice under Ex parte Quayle, 1935 C.D. 11, 453 O.G. 213. | | | | | |
| Dispositi | on of Claims | | | | | |
| 4)⊠ | Claim(s) <u>1-9,11,12,14 and 15</u> is/are pending i | n the application. | | | | |
| 4a) Of the above claim(s) is/are withdrawn from consideration. | | | | | | |
| 5) | Claim(s) is/are allowed. | | | | | |
| 6) Claim(s) is/are rejected. | | | | | | |
| 7) | 7) Claim(s) is/are objected to. | | | | | |
| 8)⊠ Claim(s) <u>1-9,11,12,14 and 15</u> are subject to restriction and/or election requirement. | | | | | | |
| Applicati | on Papers | | | | | |
| 9) 🗆 | The specification is objected to by the Examine | r. | | | | |
| 10)☐ The drawing(s) filed on is/are: a)☐ accepted or b)☐ objected to by the Examiner. | | | | | | |
| Applicant may not request that any objection to the drawing(s) be held in abeyance. See 37 CFR 1.85(a). | | | | | | |
| 11)☐ The proposed drawing correction filed on is: a)☐ approved b)☐ disapproved by the Examiner. | | | | | | |
| If approved, corrected drawings are required in reply to this Office action. | | | | | | |
| 12) The oath or declaration is objected to by the Examiner. | | | | | | |
| Priority ι | ınder 35 U.S.C. §§ 119 and 120 | | | | | |
| 13) Acknowledgment is made of a claim for foreign priority under 35 U.S.C. § 119(a)-(d) or (f). | | | | | | |
| a) ☐ All b) ☐ Some * c) ☐ None of: | | | | | | |
| | Certified copies of the priority documents have been received. | | | | | |
| | 2. Certified copies of the priority document | s have been received in A | oplication No | | | |
| 3. Copies of the certified copies of the priority documents have been received in this National Stage application from the International Bureau (PCT Rule 17.2(a)). * See the attached detailed Office action for a list of the certified copies not received. | | | | | | |
| 14) Acknowledgment is made of a claim for domestic priority under 35 U.S.C. § 119(e) (to a provisional application). | | | | | | |
| a) The translation of the foreign language provisional application has been received. 15) Acknowledgment is made of a claim for domestic priority under 35 U.S.C. §§ 120 and/or 121. | | | | | | |
| Attachmen | | and of the order | 33 Gilaror 12 1. | | | |
| 1) Notic | e of References Cited (PTO-892) e of Draftsperson's Patent Drawing Review (PTO-948) nation Disclosure Statement(s) (PTO-1449) Paper No(s) _ | 5) Notice of I | Summary (PTO-413) Paper No(s) Informal Patent Application (PTO-152) | | | |
| U.S. Patent and Ti PTO-326 (Re | _ | tion Summary | Part of Paper No. 6 | | | |

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Art Unit: 1645

1. Restriction is required under 35 U.S.C. 121 and 372.

This application contains the following inventions or groups of inventions which are not so linked as to form a single general inventive concept under PCT Rule 13.1.

In accordance with 37 CFR 1.499, applicant is required, in response to this action, to elect a single invention to which the claims must be restricted.

Election/Restrictions

- Group I Claims 1-9 and 11 are drawn to an immunogenic composition and vaccine comprising at least one *Streptococcus pneumoniae* polysaccharide antigen, at least one *Streptococcus pneumoniae* protein antigen and adjuvant. Further species election required.
- Group II Claim 12 is drawn to a method of preventing or ameliorating

 Streptoccocus pneumoniae in a patient.
- Group III Claim 14 is drawn to a method of making the immunogenic composition of claim 1.
- Group IV Claim 15 is drawn to a method of preventing or ameliorating Otitis media in infants comprising administering a safe and effective amount of a vaccine comprising a *Streptococcus pneumoniae* polysaccharide antigen, *Streptococcus pneumoniae* protein antigen and TH1 inducing adjuvant to said infant.

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2. The inventions listed as Groups I-V do not relate to a single general inventive concept under PCT Rule 13.1 because, under PCT Rule 13.2, they lack the same or corresponding special technical features for the following reasons:

Group I lacks novelty under PCT Article 33(2) as being anticipated by

Paton et al, (WO 9006951, published June 1990). Paton et al discloses a vaccine that comprises capsular polysaccharide material conjugated with an altered pneumolysin (page 3). Paton et al further disclose that the vaccine can be administered with an approved adjuvant (page 4). Group I is the main invention in this application and it lacks novelty, therefore the other claims are not so linked by a special technical feature within the meaning of PCT Rule 13.2 so as to form a single inventive concept.

3. This application contains claims directed to the following patentably distinct species of the claimed invention. In the event applicant elects Group I applicant is required to elect an antigen. Claims 1-9 and 11 are generic to plurality of disclosed patentably distinct species, based on structural and functional differences, comprising:

Species A, drawn to pneumolysin

Species B, drawn to PspA

Species C, drawn to PspC

Species D, drawn to PsaA

Species E, drawn to glyceraldehydes-3-phosphate dehydrogenase

Species F, drawn to CbpA

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Applicant is required under 35 U.S.C. 121 to elect a single disclosed species for prosecution on the merits to which the claims shall be restricted if no generic claim is finally held to be allowable.

Applicant is advised that a reply to this requirement must include an identification of the species that is elected consonant with this requirement, and a listing of all claims readable thereon, including any claims subsequently added. An argument that a claim is allowable or that all claims are generic is considered nonresponsive unless accompanied by an election.

Upon the allowance of a generic claim, applicant will be entitled to consideration of claims to additional species which are written in dependent form or otherwise include all the limitations of an allowed generic claim as provided by 37 CFR 1.141. If claims are added after the election, applicant must indicate which are readable upon the elected species. MPEP § 809.02(a).

Should applicant traverse on the ground that the species are not patentably distinct, applicant should submit evidence or identify such evidence now of record showing the species to be obvious variants or clearly admit on the record that this is the case. In either instance, if the examiner finds one of the inventions unpatentable over the prior art, the evidence or admission may be used in a rejection under 35 U.S.C. 103(a) of the other invention.

4. Species A-F are structurally independent and distinct each from the other.

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5. Applicant is advised that the reply to this requirement to be complete must include an election of invention to be examined even though the requirement be traversed (37 CFR 1.143).

6. Any inquiry of the general nature or relating to the status of this general application should be directed to the Group receptionist whose telephone number is (703) 308–0196.

Papers relating to this application may be submitted to Technology Center 1600, Group 1640 by facsimile transmission. The faxing of such papers must conform with the notice published in the Office Gazette, 1096 OG 30 (November 15, 1989). Should applicant wish to FAX a response, the current FAX number for the Group 1600 is (703) 308-4242.

Any inquiry concerning this communication from the examiner should be directed to Vanessa L. Ford, whose telephone number is (703) 308-4735. The examiner can normally be reached on Monday – Friday from 7:30 AM to 4:00 PM. If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Lynette Smith, can be reached at (703) 308–3909.

Vanessä L. Ford

Biotechnology Patent Examiner

February 10, 2002

LYNETTE R. F. SMITH
SUPERVISORY PATENT EXAMINER
TECHNOLOGY CENTER 1600